Health Certificate No. _____
Valid only if the USDA Veterinary
Seal Appears Over the Certificate No.

CHAPTER 3 (E)

Health certificate

For flavouring innards for use in the manufacture of petfood, intended for dispatch to the European Community

Note for the importer: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

1. Consignor (name and address in full) ……. ……. ……. ……. ……. …….	**VETERINARY CERTIFICATE** **For flavouring innards for use in the manufacture of petfood, intended for dispatch to the European Community** Reference number(1) ORIGINAL
2. Consignee (name and address in full) ……. ……. ……. …….	**3. Origin of the flavouring innards products** 3.1. Country :…… 3.2. Code of territory : …… **4. Competent Authority** 4.1. Responsible Ministry : …… 4.2. Certifying department : ..…….. …….
5. Destination of the flavouring innards products 5.1. EU Member State : ……..……. 5.2. Name and address of the destination : ..…….. …….. …….. …….	**6. Place of loading for exportation** ……. ……. …….. …….
7. Means of transport and consignment identification(2) 7.1. (Lorry, Rail-wagon, Ship, or Aircraft)(3) 7.2. Number of seal (if applicable) : ……..….. 7.3. Registration number(s), ship name or flight number :….. ……. …….	7.4. Nature of packaging : …….. ..……... 7.5. Number of packages : …….. 7.6. Net weight : …… 7.7. Lot/batch production reference number : ……. …….

8. Identification of the flavouring innards products

8.1. Nature of flavouring innards products :
……..……..……..

8.2. Species of animals from which theflavouring innards products derive :
…...……..……..
……..……...

8.3. Address and registration number of the approved establishment : ..……..……
……

9. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002(4) and certify that the flavouring innards products described above :

9.1. Consist of animal by-products that satisfy the animal health requirement below;

9.2. Have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;

9.3. have been prepared including the following animal by-products which are exclusively :

(3) *either* [- parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons;]

(3) *and/or* [- parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance with Community legislation;]

[3] *and/or*	[-	hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]
[3] *and/or*	[-	blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]
[3] *and/or*	[-	animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves;]
[3] *and/or*	[	former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals;]
[3] *and/or*	[-	raw milk originating from animals that do not show clinical signs of any disease communicable through that product to humans or animals,]
[3] *and/or*	[-	fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production;]
[3] *and/or*	[-	fresh by-products from fish from plants manufacturing fish products for human consumption;]
[3] *and/or*	[-	shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals;]

9.4. have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation 1774/2002/EC , in order to kill pathogenic agents;

9.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards [5]:

Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0,

Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;

9.6. the end product was :

[3] *either* [packed in new or sterilised bags,]

[3] *or* [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,]

and which bear labels indicating “NOT FOR HUMAN CONSUMPTION”;

9.7. the end product was stored in enclosed storage;

9.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.

Official stamp and signature

Done at …….. on .....
(place) (date)

(stamp) [6]

……..
(signature of the official veterinarian) [6]

……..
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.

(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.

(3) Delete as appropriate.

(4) OJ L 273, 10.10.2002, p. 1.

(5) Where:

- n = number of samples to be tested,
- m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m,
- M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more, and
- c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.

(6) The signature and the stamp must be in a different colour to that of the printing.